

K120810 FIBEROPTIC LASER DELIVERY SYSTEM

Jun 28, 2012
 104 days to decision

K120810 · Product code: **GEX** · General & Plastic Surgery
 Source: <https://www.510kdatabase.net/k120810/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 16, 2012
Decision date	Jun 28, 2012
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fiberoptic Fabrications, Inc.
Location	Wilbraham, MA, US
Contact	CAROL J MORELLO
510(k) history	1 submissions · 1 cleared · 2012-2012

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k120810/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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